URGENT: Medical Device Field Safety Corrective Action Product Recall

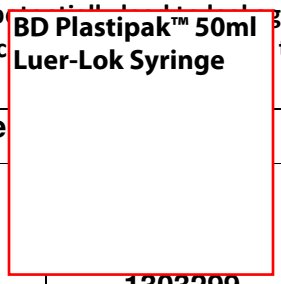
BD Reference: **MSS-13-237-FA**

Date: **26th June 2013**

Subject: **BD Plastipak™ 50ml Luer-Lok Syringe – Sterile**
Product Code: 300865:

Product Recall

BD is recalling syringes that may be potentially affected by a dent/indentation defect on the syringe barrel. This defect could p[illegible]ge throughout the stopper of the syringe. The recalled product batc[illegible] the table below.



Product Code	
300865	1303299
300865	1303300
300865	1303301
300865	1303302
300865	1303303
300865	1304202
300865	1304206
300865	1304211
300865	1304213
300865	1304217
300865	1304219
300865	1304223
300865	1304227
300865	1304228
300865	1304238

NOTE: This notification is different to the 50ml Plastipak™ Luer-Lok Syringes (Product Code 300865) recall that was initiated by BD in April 2013

YOU NEED TO TAKE THE FOLLOWING ACTIONS:

1. Please distribute this information to anyone who uses or orders **BD Plastipak™ 50ml Luer-Lok Syringes** in your institution. Additionally, please ensure that a copy of this letter is provided to any other organisations to which affected devices have been transferred.

2. Please check your inventory for BD REF 300865 with the lot numbers identified in the table above and return the product to BD.

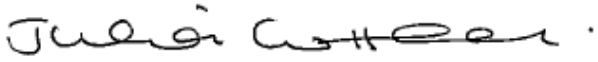
3. Please complete the Recall Response Card below and return by fax immediately to the number on the Customer Response Card.

Note: THIS FORM MUST BE COMPLETED AND FAXED TO BD MEDICAL.

4. If you have any questions regarding this communication, please contact the following telephone number 00 44 1865 781517.

Please accept our apologies for the inconvenience caused by this recall notice. We know that you share in our desire to provide superior quality products and services to both our customers and their patients.

Yours faithfully



Julie Cotterell
Marketing Manager
BD Medical – Medical Surgical Systems